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SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT BY APPLICANT

Sheet 1 of 1

Complete if Known	
Application Number	09/863,049
Filing Date	May 22, 2001
First Named Inventor	Sue J. Kenwrick
Art Unit	1632
Examiner Name	Wehbe, A.
Attorney Docket Number	HO-P01961US1

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Document Number: Number-Kind Code[2] (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear

FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Foreign Patent Document: Country Code[3]-Number[4]-Kind Code[5] (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T[6]

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NON PATENT LITERATURE DOCUMENTS

Examiner Initials*	Cite No.[1]	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T[2]
	CA**	Mayer, E J, et al; Novel corneal features in two males with incontinentia pigmenti; Br J Ophthalmol 2003; 87:554-556.	
	CB**	Bardaro, Tiziana, et al; Two Cases of Misinterpretation of Molecular Results in Incontinentia Pigmenti, and a PCR-Based Method to Discriminate NEMO/IKKy Gene Deletion; Human Mutation 21:8-11 (2002).	
	CC**	Chaturvedi, LS et al; Point mutation and polymorphism in Duchenne/Becker Muscular Dystrophy (D/BMD) patients; Experimental and Molecular Medicine, Vol. 33, No. 4,251-256, December 2001.	
	CD**	Shahbazian, Mona D. et al; Molecular genetics of Rett syndrome and clinical spectrum of MECP2 mutations; Developmental disorders 2001 Lippincott Williams & Wilkins pp. 171-176.	

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